



PTO/SB/08A (10-01)

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| <b>SUPPLEMENTAL INFORMATION<br/>DISCLOSURE<br/>STATEMENT BY APPLICANT</b> |   | Application Number       | 09/863,049      |                        |              |
|   |   | Filing Date              | May 22, 2001    |                        |              |
|   |   | First Named Inventor     | Sue J. Kenwrick |                        |              |
|   |   | Art Unit                 | 1632            |                        |              |
|   |   | Examiner Name            | Wehbe, A.       |                        |              |
| Sheet   | 1 | of                       | 1               | Attorney Docket Number | HO-P01961US1 |

| U.S. PATENT DOCUMENTS |                          |  |                                |  |  |
|-----------------------|--------------------------|--|--------------------------------|--|--|
| Examiner<br>Initials* | Cite<br>No. <sup>1</sup> | Document Number                          | Publication Date<br>MM-DD-YYYY | Name of Patentee or Applicant<br>of Cited Document | Pages, Columns, Lines,<br>Where Relevant<br>Passages or Relevant<br>Figures Appear |
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| FOREIGN PATENT DOCUMENTS |                          |   |                                |  |  |                |
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| Examiner<br>Initials*    | Cite<br>No. <sup>1</sup> | Foreign Patent Document   | Publication Date<br>MM-DD-YYYY | Name of Patentee or<br>Applicant of Cited Document | Pages, Columns, Lines,<br>Where Relevant<br>Passages or Relevant<br>Figures Appear | T <sup>6</sup> |
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|                                 |                          |  |                |
|                                 | CA**                     | Mayer, E J, et al; Novel corneal features in two males with incontinentia pigmenti; Br J Ophthalmol 2003; 87:554-556.  |                |
|                                 | CB**                     | Bardaro, Tiziana, et al; Two Cases of Misinterpretation of Molecular Results in Incontinentia Pigmenti, and a PCR-Based Method to Discriminate NEMO/IKKγ Gene Deletion; Human Mutation 21:8-11 (2002).   |                |
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|                                 | CD**                     | Shahbazian, Mona D. et al; Molecular genetics of Rett syndrome and clinical spectrum of MECP2 mutations; Developmental disorders 2001 Lippincott Williams & Wilkins pp. 171-176.   |                |

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12/4/03